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WHAT IS CLAIMED IS:

2 1. A compound of the formula:

$$R^4$$
 R^3
 R^{14}
 R^5
 R^5

4 a pharmaceutically acceptable salt or a prodrug thereof,

5 wherein

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- R¹ is alkyl or -NR⁷R⁸, where each of R⁷ and R⁸ is independently hydrogen or alkyl;
- 8 R² is hydrogen or alkyl;
- each of R³, R⁴, R⁵, and R⁶ is independently hydrogen, halide, alkyl,
- 10 –OR⁹ (where R⁹ is hydrogen, alkyl, a hydroxy protecting group,
- or cycloalkylalkyl), -SR¹⁰ (where R¹⁰ is hydrogen or alkyl), or
- $-NR^{11}R^{12}$ (where each of R^{11} and R^{12} is independently
- hydrogen, alkyl, or a nitrogen protecting group), provided R³,
- 14 R⁴, R⁵, and R⁶ are not all simultaneously alkyl); or R³ and R⁴
- together with atoms to which they are attached to form
- heterocyclyl, heteroaryl, or cycloalkyl; and
- 17 R¹⁴ is hydrogen, lower alkyl or -OR¹⁵, where R¹⁵ is hydrogen, lower
- 18 alkyl, or a hydroxy protecting group.
 - 2. The compound according to Claim 1, wherein R¹⁴ is hydrogen.
- 1 3. The compound according to Claim 2, wherein R¹ is alkyl.
- 1 4. The compound according to Claim 3, wherein R¹ is selected from the 2 group consisting of methyl, ethyl, and isopropyl.
- 1 5. The compound according to Claim 3, wherein R² is hydrogen.
- 1 6. The compound according to Claim 5, wherein each of R⁷ and R⁸ is independently hydrogen or methyl.

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1	7.	The compound according to Claim 6, wherein each of R ³ , R ⁴ , R ⁵ , and		
2	R ⁶ is independently l	nydrogen, halide, alkyl, or -OR ⁹ , where R ⁹ is hydrogen, alkyl, a hydroxy		
3	protecting group, or cycloalkylalkyl; or R ³ and R ⁴ together with atoms to which they are			
4	attached to form heterocyclyl, heteroaryl, or cycloalkyl.			
1	8.	The compound according to Claim 7, wherein at least one of R ³ , R ⁴ ,		
2	R ⁵ , and R ⁶ is alkyl, h	alide, or -OR ⁹ , where R ⁹ is as defined in Claim 1.		
1	9.	The compound according to Claim 8, wherein at least one of R ³ , R ⁴ ,		
2	R ⁵ , and R ⁶ is bromo,	chloro, fluoro, methoxy, ethoxy, methyl, and hydroxy.		
1	10.	The compound according to Claim 9, wherein		
2	(a)	R^3 is methoxy, and R^4 , R^5 , and R^6 are hydrogen;		
3	(b)	R ³ is methyl, R ⁶ is methoxy, and R ⁴ and R ⁵ are hydrogen;		
4	(c)	R ³ is methyl, R ⁶ is chloro, and R ⁴ and R ⁵ are hydrogen;		
5	(d)	R ³ is chloro, R ⁴ is methoxy, and R ⁵ and R ⁶ are hydrogen;		
6	(e)	R ³ is methyl, R ⁴ is chloro, and R ⁵ and R ⁶ are hydrogen;		
7.	(f)	R ³ is methyl, R ⁴ is methoxy, and R ⁵ and R ⁶ are hydrogen;		
8	(g)	R^4 is chloro, and R^3 , R^5 and R^6 are hydrogen;		
9	(h)	R ⁴ is methoxy, and R ³ , R ⁵ , and R ⁶ are hydrogen.		
10	(i)	R ³ is methyl, R ⁶ is bromo, and R ⁴ and R ⁵ are hydrogen;		
11	(j)	R ³ is bromo, R ⁴ is methoxy, and R ⁵ and R ⁶ are hydrogen;		
12	(k)	R ³ is methyl, R ⁴ is bromo, and R ⁵ and R ⁶ are hydrogen;		
13	(1)	R ⁴ is bromo, and R ³ , R ⁵ and R ⁶ are hydrogen; or		
14	(m)	R ³ is ethoxy and R ⁴ , R ⁵ and R ⁶ are hydrogen.		
1	11.	The compound according to Claim 7, wherein R ³ and R ⁴ together with		
2	atoms to which they are attached to form furanyl, dihydrofuranyl, or pyrrolyl.			

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with atoms to which they are attached to form furanyl or dihydrofuranyl.

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12.

The compound according to Claim 11, wherein \mathbb{R}^3 and \mathbb{R}^4 together

2 the formula:

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- 1 14. A method for producing an imidazolin-2-ylmethyl-substituted aromatic
- 2 compound of the formula:

$$\mathbb{R}^{1}$$

$$\mathbb{R}^{2}$$

$$\mathbb{R}^{2}$$

$$\mathbb{R}^{6}$$

$$\mathbb{R}^{14}$$

$$\mathbb{R}^{14}$$

$$\mathbb{R}^{14}$$

3 4

said method comprising contacting a nitrile compound of the formula:

5 6

- with ethylene diamine to produce the imidazolin-2-ylmethyl-substituted aromatic compound,
- 7 wherein
- R¹ is alkyl, -NR⁷R⁸, where each of R⁷ and R⁸ is independently hydrogen or alkyl;
- 10 R² is hydrogen or alkyl;
- each of R³, R⁴, R⁵, and R⁶ is independently hydrogen, halide, alkyl, -OR⁹,
- where R⁹ is hydrogen, alkyl, a hydroxy protecting group, or
- cycloalkylalkyl, –SR¹⁰, where R¹⁰ is hydrogen or alkyl, or –NR¹¹R¹²,
- where each of R¹¹ and R¹² is independently hydrogen, alkyl, or a
- nitrogen protecting group, provided R³, R⁴, R⁵, and R⁶ are not all
- simultaneously alkyl); or R³ and R⁴ together with atoms to which they
- are attached to form heterocyclyl, heteroaryl, or cycloalkyl; and
- 18 R¹⁴ is hydrogen, lower alkyl or -OR¹⁵, where R¹⁵ is hydrogen, lower alkyl, or
- 19 a hydroxy protecting group.

compound of the formula:

4 said method comprising contacting an ester compound of the formula:

$$R^{4}$$

$$R^{2}$$

$$R^{6}$$

$$R^{14}$$

$$R^{14}$$

$$R^{13}$$

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6 with ethylene diamine in the presence of a trialkylaluminum to produce the imidazolin-2-

ylmethyl-substituted aromatic compound,

wherein

9 R¹ is alkyl, -NR⁷R⁸, where each of R⁷ and R⁸ is independently hydrogen or alkyl;

11 R² is hydrogen or alkyl;

each of R³, R⁴, R⁵, and R⁶ is independently hydrogen, halide, alkyl, -OR⁹,

where R⁹ is hydrogen, alkyl, a hydroxy protecting group, or

cycloalkylalkyl, –SR¹⁰, where R¹⁰ is hydrogen or alkyl, or –NR¹¹R¹²,

where each of R¹¹ and R¹² is independently hydrogen, alkyl, or a

nitrogen protecting group; or R³ and R⁴ together with atoms to which

they are attached to form heterocyclyl, heteroaryl, or cycloalkyl;

18 R¹³ is alkyl; and

19 R¹⁴ is hydrogen, lower alkyl or -OR¹⁵, where R¹⁵ is hydrogen, lower alkyl, or a hydroxy protecting group.

- 1 16. The method of Claim 15, wherein the trialkylaluminum is trimethylaluminum or triethylaluminum.
- 1 17. A composition comprising:
 - (a) a therapeutically effective amount of a compound of Claim 1; and
- 3 (b) a pharmaceutically acceptable carrier.

1	1	.8.	A method for treating a disease state selected from the groups		
2	consisting of urg	ge inco	ontinence, stress incontinence, overflow incontinence, functional		
3	incontinence, sexual dysfunction, nasal congestion, and CNS disorders selected from the				
4	group depression, anxiety, dementia, senility, Alzheimer's, deficiencies in attentiveness and				
5	cognition, eating disorders, obesity, bulimia and anorexia, said method comprising				
6	administering to a patient in need of such treatment a therapeutically effective amount of a				
7	compound of Claim 1.				
1	1	.9.	A method for treating a disease state comprising urinary incontinence		
2	by administering to a subject in need of such treatment an effective amount of a Compound				
3	of Claim 1.				
4					
1	2	20.	The method of Claim 19, wherein the disorder is stress incontinence.		
1	2	21.	The method of Claim 19, wherein the disorder is urge incontinence.		
1	2	22.	A method for treating nasal congestion by administering to a mammal		
2	in need of such treatment an effective amount of a Compound of Claim 1.				
1	2	23.	The method of Claim 22, wherein the disorder is nasal congestion.		
1	2	24.	The method of Claim 23, wherein the disorder is sinusitis or otitis.		
1	. 2	25.	A method for treating sexual dysfunction by administering to a		
2	mammal in need of such treatment an effective amount of a Compound of Claim 1.				